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AMENDMENTS TO THE CLAIMS:

Set forth below in ascending order, with status identifiers, is a complete listing of all claims currently under examination. Changes to any amended claims are indicated by strikethrough and underlining. This listing also reflects any cancellation and/or addition of claims.

1. (Currently Amended) A process for separating a cell type from a mixture of cell types by

electrophoresis comprising:

(a) providing a sample comprising between about 10⁵ to about 10¹⁰ cells/mL of a mixture of cell types to a <u>first or second</u> sample chamber of <u>an</u> electrophoresis apparatus, <u>wherein said electrophoresis apparatus comprises</u> a first electrolyte chamber; a second electrolyte chamber disposed between the first electrolyte chamber and the second electrolyte chamber; a second sample chamber disposed adjacent to the first sample chamber disposed and between the first electrolyte chamber and the second electrolyte chamber; a first ion-permeable barrier disposed between the first sample chamber and the second sample chamber; a second ion-permeable barrier disposed between the first electrolyte chamber and the first sample chamber; a third ion-permeable barrier disposed between the second sample chamber and the second electrolyte chamber; and the electrodes disposed in the first and second electrolyte chambers; and

(b) applying an electric potential between the electrodes causing at least one cell type in the first sample chamber or the second sample chamber to move through the first ion-permeable barrier into the other of the first or second sample chamber.

2. (Original) The process according to claim 1 wherein the at least one cell type is selected from the group consisting of cancer, totipotent, multipotent, pluripotent, stem, viable, non-viable, bacterial, erythrocyte, leukocyte, bone marrow, organ, tissue, single cell eukaryote, prokaryote, algae, and plant.

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3. (Original) The process according to claim 2 wherein the at least one cell type is selected from

the group consisting of erythrocyte, leukocyte, bone marrow cell, organ cell, stem cell, and tissue

cell.

4. (Previously Presented) The process according to claim 1 wherein the sample contains

comprises at least two cell populations.

5. (Currently Amended) The process according to claim 1 wherein the a cell type of interest is

caused to movemoves out of the sample through the first ion-permeable barrier into the other of

the first or second sample chamber and unwanted cell types remain in the sample during

electrophoresis, or the cell type of interest remains in the sample and unwanted cell types are

caused to-move out of the sample into the other of the first or second sample chamber during

electrophoresis.

6. (Previously Presented) The process according to claim 1 wherein substantially all transbarrier

migration of a desired cell type occurs upon the application of the electric potential.

7. (Previously Presented) The process according to claim 1 wherein the first ion-permeable

barrier prevents substantial convective mixing of contents of the first and second sample

chambers, the second ion-permeable barrier prevents substantial convective mixing of contents

of the first electrolyte chamber and the first sample chamber, and the third ion-permeable barrier

prevents substantial convective mixing of contents of the second electrolyte chamber and the

second sample chamber.

8. (Previously Presented) The process according to claim 1 wherein the step of applying an

electric potential between the electrodes is maintained until at least one cell type reaches a

desired purity level in the first or second sample chamber.

9. (Previously Presented) The process according to claim 1 wherein the first ion-permeable

barrier is a membrane having a characteristic average pore size and pore size distribution.

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10. (Previously Presented) The process according to claim 1 wherein all the ion-permeable

barriers are membranes having a characteristic average pore size and pore size distribution.

11. (Previously Presented) The process according to claim 10 wherein at least a portion of the

membranes are made from polyacrylamide and have a molecular mass cut-off of at least about 5

kDa.

12. (Previously Presented) The process according to claim 10 wherein the first barrier is a large

pore sized membrane selected from the group consisting of a polycarbonate membrane, a

polyacrylamide membrane, a polyvinyl alcohol (PVA) membrane, a polyethersulfone (PES)

membrane, a polyvinylidene fluoride (PVDF) membrane, a nylon membrane, an acrylic

copolymer based membrane, a vinyl copolymer based membrane, a polysulfone membrane, a

cellulose membrane, a cellulose triacetate membrane, a cellulose ester, a polypropylene

membrane, a silicate, a borosilicate, and a glass fiber.

13. (Previously Presented) The process according to claim 12 wherein the large pore sized

membrane is a polycarbonate membrane.

14. (Previously Presented) The process according to claim 12 or 13 wherein the pore size is from

about 0.01 to about 100 µm.

15. (Previously Presented) The process according to claim 14 wherein the pore size is from about

1 to about 10 μ m.

16. (Previously Presented) The process according to claim 1 wherein the second and third

barriers are restriction membranes having a molecular mass cut off less than that of the first

barrier.

17. (Original) The process according to claim 16 wherein the restriction membranes are formed

from polyacrylamide.

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18. (Previously Presented) The process according to claim 1 wherein at least about 50% of the at

least one cell type remains viable or substantially unchanged after separation.

19. (Previously Presented) The process according to claim 18 wherein at least about 60% of the

at least one cell type remains viable or substantially unchanged after separation.

20. (Previously Presented) The process according to claim 1 wherein the sample is processed in a

static form in batches or processed in a substantially continuous form by moving the sample and

electrolyte in streams through the apparatus.

21. (Previously Presented) The process according to claim 1 wherein the difference in the

electric potential is from about 1 to about 200 V.

22. (Previously Presented) The process according to claim 21 wherein the voltage is about 60 V.

23. (Previously Presented) The process according to claim 21 wherein the field strengths are

from about 10 to about 100 V/cm.

24. (Previously Presented) The process according to claim 20 wherein the field strength is about

50 V/cm.

25. (Currently Amended) The process according to claim 1 wherein electrophoresis run times

arethe electric potential is applied for a period of from about 1 to about 60 minutes.

26. (Previously Presented) The process according to claim 25 wherein the electrophoresis run

time is about 10 minutes.

27. (Previously Presented) The process according to claim 1 wherein buffer or electrolyte

concentrations are between about 100 to about 400 mM.

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28. (Previously Presented) The process according to claim 27 wherein the buffer or electrolyte is

a cell-compatible biological buffer comprising at least one component selected from the group

consisting of HEPPS, HEPES, BisTris, sodium chloride, phosphate buffer salts, sucrose, glucose

and mannitol.

29. (Canceled)

30. (Currently Amended) The process according to claim 29-1 wherein the cell concentrations

concentration of the sample are is between about 10⁶ and about 10⁸ cells/mL.